ALLERGY- diphenhydramine hydrochloride capsule Meijer Distribution Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Meijer Distribution, Inc. Allergy Capsules Drug Facts

Active ingredient (in each capsule)

Diphenhydramine HCl 25 mg

Purpose

Antihistamine

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat
- temporarily relieves these symptoms due to the common cold:
- runny nose
- sneezing

Warnings

Do not use

- with any other product containing diphenhydramine, even one used on skin
- to make a child sleepy

Ask a doctor before use if you have

- glaucoma
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis

Ask a doctor or pharmacist before use

if you are taking sedatives or tranquilizers

When using this product

- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

• excitability may occur, especially in children

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- take every 4 to 6 hours
- do not take more than 6 doses in 24 hour

adults and children 12 years and over	1 to 2 capsules
children 6 to under 12 years	1 capsule
children under 6 years	do not use

Other information

- store at 20-25°C (68-77°F). Avoid high humidity. Protect from light.
- each capsule is sealed with a Uni Band® seal which bonds the two capsule halves together. Do not use if seal is broken or missing. Do not use if blister unit is broken or torn.

Inactive ingredients

anhydrous lactose, benzyl alcohol, butylparaben, D&C red no. 28, edetate calcium disodium, edible ink, FD&C blue no. 1, FD&C red no. 40, gelatin, magnesium stearate, methylparaben, polysorbate 80, propylparaben, sodium lauryl sulfate, sodium propionate

Questions or comments?

1-800-719-9260

Principal Display Panel

Compare to Benadryl $^{\circledR}$ active ingredient

MULTI-SYMPTOM

allergy capsules

Diphenhydramine HCl 25 mg | Antihistamine

RELIEF OF:

Sneezing; Itchy, Watery Eyes; Runny Nose & Itchy Throat

24 Capsules | actual size



OPEN OTHER END

Important: Read all product information before using. Keep this box for important information.

Drug Facts

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Uses

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■ sneezing
■ itchy, watery eyes
■ itching of the nose or throat

■ temporarily relieves these symptoms due to the common cold:

Warnings

Do not use ■ to make a child sleepy ■ with any other product containing diphenhydramine, even one used on skin

Ask a doctor before use if you have

■ a breathing problem such as emphysema or chronic bronchitis ■ glaucoma

■ trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers

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DIST. BY MEIJER DISTRIBUTION, INC. GRAND RAPIDS, MI 49544 www.meijer.com



Drug Facts (continued)

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Questions or comments? 1-800-719-9260

*This product is not manufactured or distributed by Johnson & Johnson, owner of the registered trademark Benadryl®.

PID 438861



ALLERGY

diphenhydramine hydrochloride capsule

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:41250-462

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength DIPHENHYDRAMINE HYDRO CHLO RIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - DIPHENHYDRAMINE 25 mg UNII:8GTS82S83M) HYDROCHLORIDE

Inactive Ingredients Ingredient Name Strength ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK) BENZYL ALCOHOL (UNII: LKG8494WBH) BUTYLPARABEN (UNII: 3QPI1U3FV8)

D&C RED NO. 28 (UNII: 767IP0 Y5NH)

EDETATE CALCIUM DISO DIUM (UNII: 25IH6 R4SGF)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 80 (UNII: 6 OZP39 ZG8 H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Product Characteristics			
Color	PINK (clear), WHITE (clear), RED (band)	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	L462
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-462-62	24 in 1 CARTON	0 1/15/19 9 1	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41250-462-67	48 in 1 CARTON	0 1/15/19 9 1	10/12/2016
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:41250-462-78	1 in 1 CARTON	0 1/15/19 9 1	
3		100 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	0 1/15/19 9 1	

Labeler - Meijer Distribution Inc (006959555)

Revised: 11/2020 Meijer Distribution Inc